REMARKS

The specification has been amended to include SEQ ID numbers which were omitted at the time of filing. Attached hereto is a marked-up version of the changes made to the specification by the current amendment. The attached page is captioned "Version With Markings To Show Changes Made."

The undersigned hereby states that the computer readable form copy (CRF copy) of the Sequence Listing and the paper copy of the Sequence Listing, submitted in accordance with 37 C.F.R. § 1.825(a) and (b), respectively, are the same and contain no new matter. Accordingly, entry of the Sequence Listing into the above-captioned case is respectfully requested.

In the unlikely event that the transmittal letter is separated from this document and the Patent Office determines that an extension and/or other relief is required, applicant petitions for any required relief including extensions of time and authorizes the Assistant Commissioner to charge the cost of such petitions and/or other fees due in connection with the filing of this. document to <u>Deposit Account No. 03-1952</u> referencing docket no. 300622005400. However, the Assistant Commissioner is not authorized to charge the cost of the issue fee to the Deposit Account.

Respectfully submitted,

Dated:

January 2, 2003

By:

Brenda J. Wallach, Ph.D. Registration No. (45,193)

Morrison & Foerster LLP 3811 Valley Centre Drive Suite 500

San Diego, California 92130-2332 Telephone: (858) 720-7961 Facsimile: (858) 720-5125

VERSION WITH MARKINGS TO SHOW CHANGES MADE

In the Specification:

Paragraph [58] beginning at page 21 has been amended as follows:

S. erythraea K97-71 contains a chromosomal deletion of the three eryA genes and insertion of the xylE gene from Pseudomonas aeruginosa in their place in the chromosome. To make this strain, plasmid pKOS97-49b was first constructed as follows. Two fragments flanking the eryA genes were PCR amplified from S. erythraea genomic DNA using the following primers (SphI, HindIII, BamH I, and EcoRI restriction sites are underlined): eryAI left flank, forward:

- 5'-TTT<u>GCATGC</u>GGCCACGCGCACGTCGTGACC (SEQ ID NO:1), eryAI left flank, reverse:
- 5'-TT<u>AAGCTT</u>CATATGTCCCCCTACTCGACGACCAC (SEQ ID NO:2); eryAIII right flank, forward:
- 5'-TTT<u>GGATCC</u>GGCGGAGGGAATACATGACCACGAC (SEQ ID NO:3), *eryAIII* right flank, reverse:
 - 5'-TTTGAATTCCCGCTCGCTGAAGTCCAGGCT (SEQ ID NO:4).--

Paragraph [65] beginning at page 24 has been amended as follows:

S. erythraea K24-1 contains a chromosomal deletion of the three eryA genes and insertion of the attB locus for the Streptomyces phage phiC31 from Streptomyces lividans, followed by the ermE* promoter in their place. To make this strain, plasmid pKOS134-04 was first constructed as follows. The phiC31 attB site was inserted between the Hind III and BamH I sites of pKOS97-49a using the following two annealed oligonucleotides: forward:

5'-AGCTTCGGGTGCCAGGGCGTGCCCTTGGGCTCCCCGGGCGCGTAA-CTAGTG (SEQ ID NO:5), and

reverse:

5'-GATCCACTAGTTACGCGCCCGGGGAGCCCAAGGGCACGCCCTGG-CACCCGA (SEQ ID NO:6).

This plasmid was designated pKOS024-87. Plasmid pKOS0134-04 was made by inserting a ~300 bp *NheI/BamHI* fragment containing the *ermE** promoter between the resulting *SpeI* and *BamHI* sites of pKOS024-87.